Remarks

Reconsideration of this Application is respectfully requested.

The above amendments do not add new matter. The amendments correct a formal matter without changing the scope of the claims. Specifically, the amendments are in response to the Office Communication, dated October 22, 2002, requesting the insertion of SEQ ID numbers for reference peptides found in the tables located on pages 33-179 of the specification. Applicants point out that all of the peptide sequences listed in Tables 1-3 and 5-6 are sequences contained within the full-length C35 polypeptide sequence of SEQ ID NO: 2 at the positions noted. As such, they do not require their own SEQ ID NOS. Paragraph 87 has been amended to further clarify this. Table 4, in addition to listing peptide sequences contained in SEQ ID NO: 2, also contains "modified" C35 peptide sequences containing amino acid substitutions. The modified sequences correspond to SEQ ID NOS: 85-147. Table 4 has been amended to specifically associate the SEQ ID NOS with the corresponding sequences. These changes are believed to introduce no new matter, and their entry is respectfully requested.

Conclusion

Applicants believe that a full and complete reply has been made to the outstanding Office Communication. If the Examiner believes, for any reason, that

personal communication will expedite prosecution of this application, the Examiner is invited to telephone the undersigned at the number provided.

Prompt and favorable consideration of this Amendment and Reply is respectfully requested.

Respectfully submitted,

STERNE, KESSLER, GOLDSTEIN & FOX P.L.L.C.

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SKGF_DC1:78084.1

SKGF Rev. 4/9/02

Version with markings to show changes made

-- A number of computer algorithms have been described for identification of peptides in a larger protein that may satisfy the requirements of peptide binding motifs for specific MHC class I or MHC class II molecules. Because of the extensive polymorphism of MHC molecules, different peptides will often bind to different MHC molecules. Tables 1-3 list C35 peptides predicted to be MHC binding peptides using three different algorithms. Specifically, Tables 1 and 5 list C35 HLA Class I and II epitopes predicted using the rules found at the SYFPEITHI website (wysiwyg://35/http://134.2.96.221/scripts/hlaserver.dll/EpPredict.htm) and are based on the book "MHC Ligands and Peptide Motifs" by Rammensee, H.G., Bachmann, J. and Stevanovic, S. (Chapman & Hall, New York 1997). Table 2 lists predicted MHC binding peptides derived from the C35 sequence using the NIH BIMAS program available on the web (http://bimas.dcrt.nih.gov/cgi-bin/molbio/ken parker comboform). Finally, Tables 3 and 6 list predicted C35 peptides identified by the Tepitope program, a program for prediction of peptides that may bind to multiple different MHC class II molecules. Using Tepitope, four C35 peptides were identified as likely candidates for binding to a variety of HLA class II molecules. These peptides are, in general, longer than those binding to HLA class I and more degenerate in terms of binding to multiple HLA class II molecules. Unless expressly noted otherwise, all peptide sequences listed in Tables 1-6 refer to C35 peptide sequences appearing in SEQ ID NO:2 at the amino acid positions noted. --

TABLE 4

Modifications that Enhance HLA Class I Binding

(Unless otherwise indicated, examples apply to peptides of 9 amino acids; for 10-mers the amino acid at position 5 is disregarded and the resultant 9-mer is evaluated (http://bimas.dcrt.nih.gov/cgi-bin/molbio/hla_coefficient viewing_page. The modifications listed below are provided by way of example based on current data in existing databases and are not intended in any way to be an inclusive list of all potential alterations of peptides binding all potential HLA molecules, both known and unknown to date.)

HLA A*0101

Any altered peptide that has S or T at position 2

Any altered peptide that has D or E at position 3

Any altered peptide that has P at position 4

Any altered peptide that has A, F, I, L, M, P, V, or Y at position 7

Any altered peptide that has F, K, R, or Y at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: D, E, F, G, H, K, M, N, P, Q, R, W, Y

P3: E, K, R, W

P4: K, R

P7: D, E, G, R

P9: D, E, P

HLA A*0201

Any altered peptide that has F, I, K, L, M, V, W, or Y at position 1

Any altered peptide that has I, L, M, Q, or V at anchor position 2

Any altered peptide that has F, L, M, W, or Y at position 3

Any altered peptide that has D or E at position 4

Any altered peptide that has F at position 5

Any altered peptide that has F, I, L, M, V, W or Y at auxiliary anchor position 6

Any altered peptide that has F, or W at position 7

Any altered peptide that has F, W, or Y at position 8

Any altered peptide that has I, L, T or V at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: D, E, H, P

P2: C, F, H, K, N, P, R, S, W, Y

P3: D, E, K, R

P7: D, E, G, R

P8: I, V

P9: D, E, F, G, H, K, N, P, Q, R, S, W, Y

HLA-A*0205

Any altered peptide that has F, I, K, L, M, V, W, or Y at position 1

Any altered peptide that has E, I, L, M, Q, or V at anchor position 2

Any altered peptide that has F, L, M, W, or Y at position 3

Any altered peptide that has D or E at position 4

Any altered peptide that has F, Y at position 5

Any altered peptide that has F, I, L, M, V, W or Y at auxiliary anchor position 6

Any altered peptide that has F, or W at position 7

Any altered peptide that has F, W, or Y at position 8

Any altered peptide that has I, L, T or V at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: D, E, P

P2: C, D, F, G, H, K, N, P, R, S, W, Y

P3: D, E, K, R

P7: D, E, R

P9: D, E, F, G, H, K, N, P, Q, R, S, W, Y

HLA-A*03

Any altered peptide that has G or K at position 1

Any altered peptide that has I, L, M, Q, T or V at anchor position 2

Any altered peptide that has F, I, L, M, V, W, or Y at position 3

Any altered peptide that has E, G or P at position 4

Any altered peptide that has F, I, P, V, W, Y at position 5

Any altered peptide that has F, I, L, M, or V at position 6

Any altered peptide that has F, I, L, M, W, or Y at position 7

Any altered peptide that has F, I, K, L, Q or Y at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: D, E, P

P2: D, E, F, G, H, K, N, R, S, W, Y

P7: G, K, R

P9: D, E, G, H, N, P, Q, S, T

HLA-A*1101

Any altered peptide that has G, K or R at position 1

Any altered peptide that has I, L, M, Q, T, V, Y at anchor position 2

Any altered peptide that has F, I, L, M, V, W, Y at position 3

Any altered peptide that has F, I, L, M, W or Y at position 7

Any altered peptide that has K or R at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: D, E, P

P2: D, E, G, H, K, N, R, S, W

P7: K, R

P9: C, D, E, G, N, P, Q, S, T

HLA-A24

Any altered peptide that has K or R at position 1

Any altered peptide that has F or Y at anchor position 2

Any altered peptide that has E, I, L, M, N, P, Q, or Vat position 3

Any altered peptide that has D, E, or P at position 4

Any altered peptide that has I, L, or V at position 5

Any altered peptide that has F at position 6

Any altered peptide that has N or Q at position 7

Any altered peptide that has E or K at position 8

Any altered peptide that has F, I, L, or M at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: D, E, H, K, R

P9: D, E, G, H, K, P, Q, R

HLA-A*3101

Any altered peptide that has K or R at position 1

Any altered peptide that has F, I, L, M, Q, T, V, or Y at anchor position 2

Any altered peptide that has F, I, L, M, V W, or Y at position 3

Any altered peptide that has F, I, L, M, or V at position 6

Any altered peptide that has F, I, L, M, W, or Y at position 7

Any altered peptide that has K or R at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: D, E, P

P2: D, E, G, H, K, N, R, S

P7: K, R

P9: C, G, N, P, Q, S, T

HLA-A*3302

Any altered peptide that has D or E at position 1

Any altered peptide that has I, L, M, S, V or Y at anchor position 2

Any altered peptide that has R at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: K, P, R

P2: D, E, K, R

P9: D, E, F, G, N, P, W, Y

HLA-B7

Any altered peptide that has A at position 1

Any altered peptide that has A, P or V at anchor position 2

Any altered peptide that has M or R at position 3

Any altered peptide that has P at position 5

Any altered peptide that has R at position 6

Any altered peptide that has I, L, M or V at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: D, E, F, H, K, R, W, Y

P3: D, E

P9: D, E, F, G, H, K, N, P, Q, R, S, W, Y

HLA-B8

Any altered peptide that has D or E at position 1

Any altered peptide that has A, C, L, or P at anchor position 2

Any altered peptide that has K or R at position 3

Any altered peptide that has D or E at position 4

Any altered peptide that has K or R at position 5

Any altered peptide that has I, L, M, or V at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: K, P, R

P2: D, E, F, G, H, K, Q, R, W, or Y

P3: D, E

P5: D, E

P9: D, E, F, G, H, K, N, P, Q, R, S, W, Y

HLA-B8 (8-mer peptides)

Any altered peptide that has D or E at position 1

Any altered peptide that has A, C, L, or P at anchor position 2

Any altered peptide that has K or R at position 3

Any altered peptide that has D or E at position 4

Any altered peptide that has K or R at position 5

Any altered peptide that has I, L, M, or V at anchor position 8

Any altered peptide where deleterious residues at the following positions are replaced:

P1: K, P, R

P2: D, E, F, G, H, K, Q, R, W, or Y

P3: D, E

P5: D, E

P8: D, E, F, G, H, K, N, P, Q, R, S, W, Y

HLA-B14

Any altered peptide that has D or E at position 1

Any altered peptide that has K or R at anchor position 2

Any altered peptide that has F, I, L, M, P, V, W, Y at position 3

Any altered peptide that has H or R at position 5

Any altered peptide that has I, L, M, R, or V at position 6

Any altered peptide that has T at position 7

Any altered peptide that has I, L, M, or V at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: D, E, F, W, or Y

P3: E, R

P5: E, W, Y

P9: D, E, G, H, K, N, P, Q, R

HLA-B*2702

Any altered peptide that has K or R at position 1

Any altered peptide that has E, L, M, N, Q or R at anchor position 2

Any altered peptide that has F, W, or Y at position 3

Any altered peptide that has F, I, L, W or Y at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: D, E, P

P2: D, F, G, H, K, W, or Y

P7: K

P9: D, E, G, K, N, P, Q, R, S

HLA-B27*05 (8-mer peptides)

Any altered peptide that has K or R at position 1

Any altered peptide that has E, L, M, N, Q or R at anchor position 2

Any altered peptide that has F, W, or Y at position 3

Any altered peptide that has F, I, K, L, M, R, V or Y at anchor position 8

Any altered peptide where deleterious residues at the following positions are replaced:

P1: D, E, P

P2: D, F, G, H, K, W, or Y

P7: K

P9: D, E, G, K, N, P, Q, R, S

HLA-B*3501 (8-mer peptides)

Any altered peptide that has K or R at position 1

Any altered peptide that has A, P, or S at anchor position 2

Any altered peptide that has K or R at position 3

Any altered peptide that has D or E at position 4

Any altered peptide that has D or E at position 5

Any altered peptide that has F, I, L, M, V, W or Y at anchor position 8

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: D, E, F, H, K, R, W, Y

P3: D, E

P8: D, E, F, G, H, K, P, Q, R

HLA-B*3701

Any altered peptide that has D or E at anchor position 2

Any altered peptide that has I or V at position 5

Any altered peptide that has F, L, or M at position 8

Any altered peptide that has F, I, L, M, V or Y at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P9: D, E, G, H, K, P, Q, R

HLA-B*3801

Any altered peptide that has F, H, P, W or Y at anchor position 2

Any altered peptide that has D or E at position 3

Any altered peptide that has D, E, or G at position 4

Any altered peptide that has A, I, L, M, or V at position 5

Any altered peptide that has K or Y at position 8

Any altered peptide that has F, I, L, M, or V at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

- P1: P
- P2: D, E, K, R
- P3: K, R
- P9: D, E, G, H, K, P, Q, R

HLA-B*3901 (8-mer peptides)

Any altered peptide that has H or R at anchor position 2

Any altered peptide that has D, E, F, I, L, M, V, W, or W at position 3

Any altered peptide that has D or E at position 4

Any altered peptide that has I, L, M, or V at position 6

Any altered peptide that has I, L, M or V at anchor position 8

Any altered peptide where deleterious residues at the following positions are replaced:

- P1: P
- P2: D, E
- P3: K, R
- P6: D, E, K, R
- P8: D, E, G, H, K, P, Q, R

HLA-B*3902

Any altered peptide that has K or Q at anchor position 2

Any altered peptide that has F, I, L, M, V, W, or Y at position 5

Any altered peptide that has F, L, or M at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

- P1: P
- P2: D, E
- P3: K, R
- P9: D, E, G, H, K, P, Q, R

HLA-B40

Any altered peptide that has A or G at position 1

Any altered peptide that has D or E at anchor position 2

Any altered peptide that has A, F, I, L, M, V, W, or Y at position 3

Any altered peptide that has P at position 4

Any altered peptide that has P at position 5

Any altered peptide that has A, L, M, or W at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: F, H, I, K, L, M, Q, R, V, W, or Y

P3: D, E, K, R

P9: D, E, G, H, K, N, P, Q, R

HLA-B44*03

Any altered peptide that has A, D, or S at position 1

Any altered peptide that has D or E at anchor position 2

Any altered peptide that has A, I, L, M, or V at position 3

Any altered peptide that has F, I, or P at position 4

Any altered peptide that has A, K, or V at position 5

Any altered peptide that has A, L, T, or V at position 6

Any altered peptide that has F, K, or T at position 7

Any altered peptide that has K at position 8

Any altered peptide that has F, W or Y at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: F, H, I, K, L, M, Q, R, V, W, Y

P9: D, E, G, H, K, N, P, Q, R

HLA-B*5101 (8-mer peptides)

Any altered peptide that has D, E, F, I, L, M, V, or Y at position 1

Any altered peptide that has A, G or P at anchor position 2

Any altered peptide that has F, W or Y at position 3

Any altered peptide that has D, E, G, I, K, or V at position 4

Any altered peptide that has A, G, I, S, T, or V at position 5

Any altered peptide that has I, K, L, N, or Q at position 6

Any altered peptide that has D, K, Q, or R at position 7

Any altered peptide that has I, L, M, or V at anchor position 8

Any altered peptide where deleterious residues at the following positions are replaced:

P1: K, P, R P2: D, E, H, K

P8: D, E, F, G, H, K, N, P, Q, R, S, W, Y

HLA-B*5102

Any altered peptide that has F or Y at position 1

Any altered peptide that has A, G, or P at anchor position 2

Any altered peptide that has F, I, L, V, W, or Y at position 3

Any altered peptide that has E, G, H, K, L, N, Q, R, or T at position 4

Any altered peptide that has G, N, Q, T, or V at position 5

Any altered peptide that has I, N, Q, or T at position 6

Any altered peptide that has E, K, Q, or R at position 7

Any altered peptide that has K, R, T, or Y at position 8

Any altered peptide that has I, L, M, or V at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: D, E, H, K, R

P3: D, E, K, R

P9: D, E, F, G, H, K, N, P, Q, R, S, W, Y

HLA-B*5102 (8-mer peptides)

Any altered peptide that has F or Y at position 1

Any altered peptide that has A, G, or P at anchor position 2

Any altered peptide that has F, I, L, V, W, or Y at position 3

Any altered peptide that has E, G, H, K, L, V, W, or Y at position 4

Any altered peptide that has G, N, Q, T, V at position 5

Any altered peptide that has I, N, or Q at position 6

Any altered peptide that has Q, or R at position 7

Any altered peptide that has I, L, M, or V at position 8

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: D, E, H, K, R

P3: D, E, K, R

P8: D, E, F, G, H, K, N, P, Q, R, S, W, Y

HLA-B*5103

Any altered peptide that has D, T, or V at position 1

Any altered peptide that has A, G, or P at anchor position 2

Any altered peptide that has D, F, L, or Y at position 3

Any altered peptide that has E, G, L, N, Q, R, T, or V at position 4

Any altered peptide that has A, G, M, N, Q, R, K or V at position 5

Any altered peptide that has I, K, or T at position 6

Any altered peptide that has M or V at position 7

Any altered peptide that has I, L, M, or V at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: D, E, H, K, R

P9: D, E, F, G, H, K, N, P, Q, R, S, W, Y

HLA-B*5201 (8-mer peptides)

Any altered peptide that has I, L, M, or V at position 1

Any altered peptide that has G, P, or Q at anchor position 2

Any altered peptide that has D, F, I, L, P, W, or Y at position 3

Any altered peptide that has A, E, I, K, L, P, or V at position 4

Any altered peptide that has A, F, G, I, L, M, T or V at position 5

Any altered peptide that has K, L, N, S or T at position 6

Any altered peptide that has E, K, Q, or Y at position 7

Any altered peptide that has F, I, L, M, or V at anchor position 8

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: H, K, R

P3: R

P8: D, E, G, H, K, N, P, Q, R, S

HLA-B*5801

Any altered peptide that has I, K, or R at position 1

Any altered peptide that has A, S, or T at anchor position 2

Any altered peptide that has D at position 3

Any altered peptide that has E, K, or P at position 4

Any altered peptide that has F, I, L, M, or V at position 5

Any altered peptide that has F, I, L, or V at position 6

Any altered peptide that has L, M, N, or Y at position 7

Any altered peptide that has K, N, R, or T at position 8

Any altered peptide that has F, W, or Y at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: D, E, P

P2: D, E, F, H, I, K, L, M, N, Q, R, V, W, Y

P9: D, E, G, H, K, N, P, Q, R, S

HLA-B*60

Any altered peptide that has D or E at anchor position 2

Any altered peptide that has A, I, L, M, S, or V at position 3

Any altered peptide that has L, I, or V at position 5

Any altered peptide that has I, L, M, V, or Y at position 7

Any altered peptide that has K, Q, or R at position 8

Any altered peptide that has I, L, M, or V at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: F, H, I, K, L, M, Q, R, V, W, Y

P9: D, E, F, G, H, K, N, P, Q, R, S, W, Y

HLA-B*61

Any altered peptide that has G or R at position 1

Any altered peptide that has D or E at anchor position 2

Any altered peptide that has A, F, I, L, M, T, V, W, or Y at position 3

Any altered peptide that has I at position 6

Any altered peptide that has Y at position 7

Any altered peptide that has A, I, L, M, or V at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: F, H, I, K, L, M, Q, R, V, W, Y

P9: D, E, F, G, H, K, N, P, Q, R, S, W, Y

HLA-B*61 (8-mer peptides)

Any altered peptide that has G or R at position 1

Any altered peptide that has D or E at anchor position 2

Any altered peptide that has A, F, I, L, M, T, V, W, or Y at position 3

Any altered peptide that has I at position 6

Any altered peptide that has Y at position 7

Any altered peptide that has A, I, L, M, or V at anchor position 8

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: F, H, I, K, L, M, Q, R, V, W, Y

P8: D, E, F, G, H, K, N, P, Q, R, S, W, Y

HLA-B*62

Any altered peptide that has I at position 1

Any altered peptide that has I, L, Q at anchor position 2

Any altered peptide that has G, K, R at position 3

Any altered peptide that has D, E, G, or P at position 4

Any altered peptide that has F, G, I, L, or V at position 5

Any altered peptide that has I, L, T, V at position 6

Any altered peptide that has T, V, or Y at position 7

Any altered peptide that has F, W, Y at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: D, E, F, H, K, N, R, S, W, Y

P3: D, E

P6: D, E, K, R

P9: D, E, G, H, K, N, P, Q, R, S

HLA-Cw0301

Any altered peptide that has A or R at anchor position 2

Any altered peptide that has F, I, L, M, V, or Y at position 3

Any altered peptide that has E, P, or R at position 4

Any altered peptide that has N at position 5

Any altered peptide that has F, M, or Y at position 6

Any altered peptide that has K, M, R, or S at position 7

Any altered peptide that has T at position 8

Any altered peptide that has F, I, L, M at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P3: D, K, R

P6: D, E, K, R

P9: D, E, G, H, K, N, P, Q, R, S,

HLA-Cw0401

Any altered peptide that has F, P, W, or Y at anchor position 2

Any altered peptide that has D, or H at position 3

Any altered peptide that has D or E at position 4

Any altered peptide that has A, H, M, R, or T at position 5

Any altered peptide that has I, L, M, or V at position 6

Any altered peptide that has A at position 7

Any altered peptide that has H, K, or S at position 8

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Examples of predicted human Class I MHC binding peptides - continued Subsequence

Rank Start Position

(estimated half time of dissociation)

SEQ ID NO.

Any altered peptide that has F, I, L, M, V or Y at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P2: D, E, H, K, R

P9: D, E, G, H, K, N, P, Q, R, S

HLA-Cw0602

Any altered peptide that has F, I, K, or Y at position 1

Any altered peptide that has A, P, Q, or R at anchor position 2

Any altered peptide that has F, I, K, L, or M at position 5

Any altered peptide that has I, L, or V at position 6

Any altered peptide that has K, N, Q, or R at position 7

Any altered peptide that has I, L, M, V, or Y at anchor position 9

Any altered peptide where deleterious residues at the following positions are replaced:

P1: P

P9: D, E, G, H, K, N, P, Q, R, S

Examples of predicted human Class I MHC binding peptides from the C35 aa sequence and how they might be changed to improve binding:

HLA-A*0101

Rank	Start Position	Subsequence	Score (estimated h	nalf time of dissociation)	SEQ ID NO.
1	77	KLENGGRPY	225.000		
2	16	EVEPGSGVR	90.000		
3	29	YCEPCGFEA	45.000		
4	39	YLELASAVK	36.000		
5	2	S <i>G</i> EPGQTSV	2.250	G is deleterious at P2	
exam _j	ple of ved peptide	STEPGQTSV	22.50	G replaced with T @ P2	SEQ ID NO:85

			MHC bindir	ng peptides – continued	Appl. No. 09/824,787
Rank	Start Position	Subsequence	Score (estimated h	alf time of dissociation)	SEQ ID NO.
examţ impro	ole of ved peptide	STEPGQISY	5625.00	V at P9 replaced with Y, P7 enhanced	SEQ ID NO:86
HLA-	·A*0101 (10-	-mer peptides)			
1	66	EIEINGQLVF	45.000		
2	16	EVEPGSGVRI	18.000		
3	29	YCEPCGFEAT	9.000		
4	26	VVEYCEPCGF	9.000		
5	52	GIEIESRLGG	2.250		
examţ impro	ole of ved peptide	GTEPSRLGY	1125.000	replace I with T @P2 replace G with Y @P9 P5 enhanced with P	SEQ ID NO:87
HLA-	·A*0201 (9-	mer peptides)			
1	9	SVAPPPEEV	2.982		
2	104	KITNSRPPC	2.391		
3	105	ITNSRPPCV	1.642		
4	25	IVVEYCEPC	1.485		
5	65	FEIEINGQL	1.018		
examţ impro		FLIEINWYL	16619.000		SEQ ID NO:88
HLA-	·A*0201 (10·	-mer peptides)			
1	58	RLGGTGAFEI	60.510		
2	104	KITNSRPPCV	33.472		
3	65	FEIEINGQLV	25.506		
4	83	F <i>P</i> YEKDLIEA	4.502	P is deleterious at P2	
examp impro	ole of ved peptide	FLYEKDLIEA	689.606	replace P with L @ P2	SEQ ID NO:89

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	mples of predick Start Position	cted human Class	Score	ng peptides - continued half time of dissociation)	SEQ ID NO.
	mple of roved peptide	FLYEKDLIEV	9654.485	replace A with V @ P9	SEQ ID NO:90
5	33	CGFEATYLEL	3.173		
HL	A-A*0205				
1	65	FEIEINGQL	8.820		
2	25	IVVEYCEPC	3.060		
3	9	SVAPPPEEV	2.000		
4	104	KITNSRPPC	1.500		
5	81	G <i>G</i> FPYEKDL	1.260	G is deleterious at P2	
	nple of roved peptide	GVFPYEKDL	50.400	replace G with V @ P2	SEQ ID NO:91
HL	A-A*0205 (10	-mer peptides)			
1	33	CGEFATYLEL	6.300	G is deleterious at P2	
	nple of roved peptide	CVEFATYLEL	11.200	replace G with V @ P2	SEQ ID NO:92
2	104	KITNSRPPCV	6.000		
3	65	FEIEINGQLV	2.520		
4	53	IEIESRLGGT	1.428		
5	83	FPYEKDLIEA	1.350	P is deleterious at P2	
	nple of roved peptide	FVYEKDLIEA	54.000	replace P with V @ P2	SEQ ID NO:93
HL	A-A24				
1	34	GFEATYLEL	33.000		
2	49	QYPGIEIES	11.550		
	mple of roved peptide	QYPGIEIEL	462.000	enhance P9	SEQ ID NO:94
3	70	NGQLZFSKL	11.088		

Exan	nples of predic	cted human Class	I MHC bindi	ng peptides – continued	Appl. No. 09/824,/8/
Rank	Start Position	Subsequence	Score (estimated l	half time of dissociation)	SEQ ID NO.
4	38	TYLELASAV	10.800		
5	82	GFPYEKDLI	7.500		
HLA	-A24 (10-me	r peptides)	•		
1	64	AFEIEINGQL	42.000		
2	74	VFSKLENGGF	10.000		
3	84	PYEKDLIEAI	9.000		
4	69	INGQLVFSKL	7.392		
	ple of oved peptide	IYGQLVFSKL	369.6	enhance P2	SEQ ID NO:95
5	28	EYCEPCGFEA	6.600		
HLA	A-A3				
1	77	KLENGGFPY	36.000		
	aple of oved peptide	KLENGGFPK	180.000	enhance P9	SEQ ID NO:96
2	39	YLELASAVK	20.000		
3	101	TLEKITNSR	6.000		
4	61 .	GTGAFEIEI	0.540		
5	69	INGQLVFSK	0.360	N is deleterious @ P2	
	nple of oved peptide	ILGQLVFSK	180.000	replace N with L @ P2	SEQ ID NO:97
HLA	A-A3 (10-mer	peptides)			
1	68	EINGQLVFSK	8.100		
2	58	RLGGTGAFEI	2.700		
3	41	ELASAVKEQY	1.800		
4	78	L <i>E</i> NGGFPYEK	0.810	E is deleterious @ P2	
	nple of oved peptide	LLNGGFPYEK	270.000	replace E with L @ P2	SEQ ID NO:98

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					A A
				ling peptides – c ntinued	
Kanı	Start Position	Subsequence	Score (estimated	l half time of dissociation)	SEQ ID NO.
	95	D A CNICETI EV	•	than time of dissociation)	DEQ ID NO
5	93	RASNGETLEK	0.400		
HLA	A- A*1101				
1	39	YLELASAVK	0.400		
2	69	INGQLVFSK	0.120	N is deleterious @ P2	
	nple of oved peptide	IVGQLVFSK	6.000	replace N with V @ P2	SEQ ID NO:99
3	16	EVEPGSGVR	0.120		
4	101	TLEKITNSR	0.080		
5	61	GTGAFEIEI	0.060		
HLA	A-A*1101 (10	-mer peptides)			
1	95	RASNGETLEK	1.200		
2	38	TYLELASAVK	0.600		
3	68	EINGGLVFSK	0.360		
4	78	L <i>E</i> NGGFPYEK	0.120	E is deleterious @ P2	
	nple of oved peptide	LVNGGFPYEK	4.000	replace E with V @ P2	SEQ ID NO:100
5	100	ETLEKITNSR	0.090		
HLA	A-A*3101				
1	101	TLEKITNSR	2.000		
2	16	EVEPGSGVR	0.600		
3	50	YPGIEIESR	0.400		
4	87	K <i>D</i> LIEAIRR	0.240	D is deleterious @ P2	
	nple of oved peptide	KILIEAIRR	12.000	replace D with I @ P2	SEQ ID NO:101
5	39	YLELASAVK	0.200		

					Appl. 140. 07/024,707
	nples of predi Start Position	cted human Class I Subsequence	Score	ing peptides – c ntinued half time of dissociation)	SEQ ID NO.
——— НІ. 4	A-A*3302		Commica	man time of dissociation)	
1	16	EVEPGSGVR	45.000		
2	101	TLEKITNSR	9.000		
3	50	YPGIEIESR	3.000		
4	66	EIEINGQLV	1.500		
5	56	ESRLGGTGA	1.500		
HLA	A-A*3302 (10	-mer peptides)			
1	49	QYPGIEIESR	15.000		
2	100	ETLEKITNSR	9.000		
3	16	EVEPGSGVRI	1.500		
4	28	EYCEPCGFEA	1.500		
5	68	EINGQLVFSK	1.500		
HLA	A-A68.1				
1	16	EVEPGSGVR	900.000		
2	9	SVAPPPEEV	12.000		
3	50	YPGIEIESR	10.000		·
	nple of oved peptide	YVGIEIESR	400.000	enhance P2	SEQ ID NO:102
4	96	ASNGETLEK	9.000		
5	101	TLEKITNSR	5.000		
HLA	A-A68.1 (10-n	ner peptides)			
1	100	ETLEKITNSR	300.000		
2	16	EVEPGSGVRI	18.000		
3	68	EINGGLVFSK	9.000		
4	15	E <i>E</i> VEPGSGVR	9.000	E is deleterious @ P2	

_					Appl. No. 09/824,787
	nples of predic	cted human Class I Subsequence	MHC bindir Score	ng peptides – continued	
	Position			alf time of dissociation)	SEQ ID NO.
exan	ple of				
impr	oved peptide	EVVEPGSGR	1200.00	replace E with V @ P2	SEQ ID NO:103
5	95	RASNGETLEK	3.000		
HLA	\-B14				
1	94	RRASNGETL	20.000		
2	57	SRLGGTGAF	5.000		
	ple of				
impr	oved peptide	SRLGGTGAL	100.000	enhance P9	<u>SEQ ID NO:104</u>
3	100	ETLEKITNS	3.375		
4	105	ITNSRPPCV	2.000		
5	88	DLIEAIRRA	1.350		
HLA	-B14 (10- me	r peptides)			
1	103	EKITNSRPPC	6.750		
exan	ple of				
impr	oved peptide	ERITNSRPPL	900.000	enhance P10	SEQ ID NO:105
2	33	CGFEATYLEL	5.000		
3	93	IRRASNGETL	4.000		
4	18	EPGSGVRIVV	3.000		
5	88	DLIEAIRRAS	2.250		
TTT /	D40				
HLA	A-B40				
1	65	FEIEINGQL	80.000		
2	3	GEPGQTSVA	40.000		
3	35	FEATYLELA	40.000		
4	15	EEVEPGSGV	24.000		
	aple of oved peptide	EEVEPGSGL	120.000	enhance P9	SEQ ID NO:106
5	67	IEINGQLVF	16.000		

Examples of predicted human Class I MHC binding peptides - continued

Ranl	k Start Position	Subsequence	Score (estimated l	half time of dissociation)	SEQ ID NO.
HLA	A-B40 (10-me	r peptides)			
1	55	IESRLGGTGA	20.000		
2	53	IEIESRLGGT	16.000		
	nple of oved peptide	IEIESRLGGL	80.000	enhance P10	SEQ ID NO:107
3	65	FEIEINGQLV	16.000		
4	67	IEINGQLVFS	16.000		
5	99	GETLEKITNS	8.000		
HLA	A-B60				
1	65	FEIEFNGQL	387.200		
2	17	VEPGSGVRI	17.600		
	nple of oved peptide	VEPGSGVRL	352.000	enhance P9	SEQ ID NO:108
3	15	EEVEPGSGV	16.000		
4	47	KEQYPGIEI	16.000	·	
5	85	YEKDLIEAI	8.800		
HLA	A-B60 (10-me	r peptides)			
1	65	FEIEINGQLV	16.000		
	nple of roved peptide	FEIEINGQLL	320.000	enhance P10	SEQ ID NO:109
2	106	TNSRPPCVIL	16.000		
3	53	IEIESRLGGT	8.000		
4	33	CGFEATYLEL	8.000		
5	17	VEPGSGVRIV	8.000		

	Start	Subsequence	Score	ng peptides – continued	
	Position			nalf time of dissociation)	SEQ ID NO.
HLA	B61				
1	15	EEVEPGSGV	80.000		
2	35	FEATYLELA	40.000		
exam	ple of				
	oved peptide	FEATYLELV	160.000	enhance P9	SEQ ID NO:110
3	3	GEPGQTSVA	22.000		
4	65	FEIEINGQL	16.000		
5	85	YEKDLIEAI	16.000		
HLA	-B61 (10-me	r peptides)			
1	65	FEIEINGQLV	80.000		
2	17	VEPGSGVRIV	40.000		
3	55	IESRLGGTGA	20.000		
4	87	KDLIEAIRRA	10.000		
	ple of oved peptide	KELIEAIRRV	160.000	enhance P2, P10	SEQ ID NO:111
5	53	IEIESRLGGT	8.000		
HLA	B62				
1	77	KLENGGFPY	24.000		
2	21	SGVRIVVEY	4.800		
3	75	FSKLENGGF	3.000		
4	31	EPCGFEATY	2.640	P is deleterious @ P2	
	ple of oved peptide	E Q CGFEATY	105.6	replace P with Q @ P2	SEQ ID NO:112
5	88	DLIEAIRRA	2.200		

					Appl. 140. 07/024,707
	ples of predic Start Position	cted human Class I Subsequence	Score	g peptides – c ntinued alf time of dissociation)	SEQ ID NO.
HLA	-B62 (10-me	r peptides)			
1	41	ELASAVKEQY	40.000		
2	58	RLGGTGAFEI	9.600		
3	66	EIEINGQLVF	7.920		
4	56	ESRLGGTGAF	6.000	S is deleterious @ P2	
	ple of oved peptide	EQ RLGGTGAF	480.000	replace S with Q @ P2	SEQ ID NO:113
5	20	GSGVRIVVEY	4.800	S is deleterious @ P2	
	ple of oved peptide	G Q GVRIVVEY	384.000	replace S with Q @P2	SEQ ID NO:114
HLA	-B7				
1	107	NSRPPCVIL	60.000		
	ple of oved peptide	NPRPPCVIL	1200.000	enhance P2	SEQ ID NO:115
2	45	AVKEQYPGI	6.000		
3	22	GVRIVVEYC	5.000		
4	70	NGQLVFSKL	4.000		
5	81	GGFPYEKDL	4.000		
HLA	-B7 (10-mer	peptides)			
1	50	YPGIEIESRL	80.000	•	
2	31	EPCGFEATYL	80.000		
3	18	EPGSGVRIVV	6.000		
	ple of oved peptide	EPGSGVRIVL	120.000	enhance P10	SEQ ID NO:116
4	106	TNSRPPCVIL	6.000		
5	80	NGGFPYEKDL	4.000		

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Exa	mples of predic	cted human Class	I MHC bind	ling peptides – continued	Appl. No. 09/824,/8/
	k Start Position	Subsequence	Score	half time of dissociation)	SEQ ID NO.
HL	A-B8				
1	107	NSRPPCVIL	4.000		
2	45	AVKEQYPGI	1.500		
3	105	ITNSRPPCV	0.600		
4	56	ESRLGGTGA	0.400		
5	100	ETLEKITNS	0.300	S is deleterious @ P9	
	mple of roved peptide	ETLEKITNL	12.000	replace S with L @ P9	SEQ ID NO:117
HL	A-B8 (8-mer p	peptides)			
1	83	FPYEKDLI	6.000		
2	107	NSRPPCVI	1.000		
3	91	EAIRRAS <i>N</i>	0.800	N is deleterious @ P8	
	mple of roved peptide	EAIRRASL	32.000	replace N with L @ P9	SEQ ID NO:118
4	20	GSGVRIVV	0.600		
5	18	EPGSGVRI	0.400		
HL.	A-B8 (10-mer	peptides)			
1	50	YPGIEIESRL	0.800		
2	93	IRRASNGETL	0.400		
	mple of roved peptide	IA RASNGETL	16.000	replace R with A @ P2	SEQ ID NO:119
3	31	EPCGFEATYL	0.320		
4	104	KITNSRPPCV	0.300		
5	18	EPGSGVRIVV	0.240		

	ples of predic Start Position	cted human Class I Subsequence	Score	ng peptides – c ntinued nalf time of dissociation)	SEQ ID NO.
		-	(cstillated i	ian time of dissociation)	SEQ ID NO.
HLA	-B*2702				
1	57	SRLGGTGAF	200.000		
2	94	RRASNGETL	180.000		
	ple of oved peptide	RRASNGETF	600.000	enhance P9	SEQ ID NO:120
3	93	IRRASNGET	20.000		
4	27	VEYCEPCGF	15.000		
5	77	KLENGGFPY	9.000		
HLA	-B*2702 (10-	-mer peptides)			
1	93	IRRASNGETL	60.000		
2	94	RRASNGETLE	6.000		
3	30	CEPCGFEATY	3.000		
4	58	RLGGTGAFEI	2.700		
5	23	VRIVVEYCE <i>P</i>	2.000	P is deleterious @ P10	
	ple of oved peptide	VRIVVEYCEY	200.000	replace P with Y @ P10	SEQ ID NO:121
HLA	-B*2705				
1	94	RRASNGETL	6000.000		
2	57	SRLGGTGAF	1000.000		
3	93	IRRASNGET	200.000		
	ple of oved peptide	IRRASNGEL	2000.000	enhance P9	SEQ ID NO:122
4 .	27	VEYCEPCGF	75.000		
5	77	KLENGGFPY	45.000		

Exam	ples of predic	cted human Class I	MHC bindir	ng peptides – continued	Appl. No. 09/824,787
Rank	Start Position	Subsequence	Score (estimated h	alf time of dissociation)	SEQ ID NO.
HLA	-B*2705 (10-	mer peptides)			
1	93	IRRASNGETL	2000.000		
2	94	RRASNGETL <i>E</i>	60.000	E is deleterious @ P2	
	ple of oved peptide	RRASNGETLL	6000.000	replace E with L @ P2	SEQ ID NO:123
3	78	LENGGFPYEK	30.000		
4	95	RASNGETLEK	30.000		
5	58	RLGGTGAFEI	27.000		
HLA	-B*3501				
1	31	EPCGFEATY	40.000		
2	75	FSKLENGGF	22.500		
	ple of oved peptide	FPKLENGGM	120.000	enhance P2, P9	SEQ ID NO:124
3	107	NSRPPCVIL	15.000		
4	42	LASAVKEQY	6.000		
5	18	EPGSGVRIV	4.000		
HLA	-B*3501 (10-	-mer peptides)			
1	31	EPCGFEATYL	30.000		
2	50	YPGIEIESRL	20.000		
3	56	ESRLGGTGAF	15.000		
4	20	GSGVRIVVEY	10.000		
5	83	FPYEKDLIEA	6.000		
	ple of oved peptide	FPYEKDLIEM	120.000	enhance P10	SEQ ID NO:125

				ing peptides – continued	Appl. No. 09/824,787
Rank	Start Position	Subsequence	Score (estimated)	half time of dissociation)	SEQ ID NO.
HLA	-B*3701				
1	65	FEIEINGQL	15.000		
	ple of oved peptide	F D IEINGQL	60.000	enhance P2	SEQ ID NO:126
2	47	KEQYPGIEI	10.000		
3	85	YEKDLIEAI	10.000		
4	17	VEPGSGVRI	10.000		
5	35	FEATYLELA	5.000		
HLA	-B*3701 (10-	-mer peptides)			
1	65	FEIEINGQLV	10.000		
	ple of oved peptide	FDIEINGQLI	200.000	enhance P2, P10	SEQ ID NO:127
2	67	IEINGQLVFS	5.000		
3	81	GGFPYEKDLI	5.000		
4	87	KDLIEAIRRA	4.000		
5	30	CEPCGFEATY	2.000		
HLA	-B*3801				
1	34	GFEATYLEL	6.000		
	ple of oved peptide	GHEATYLEL	90.000	enhance P2	SEQ ID NO:128
2	70	NGQLVFSKL	1.560		
3	38	TYLELASAV	1.040		
4	81	GGFPYEKDL	1.000		
5	97	SNGETLEKI	0.720		

				ing peptides – continued	Appl. No. 09/824,/8/
Rank	Start Position	Subsequence	Score (estimated	half time of dissociation)	SEQ ID NO
HLA-	-B*3801 (10-	-mer peptides)			
1	64	AFEIEINGQL	7.800		
exam _l impro	ple of wed peptide	AHEIEINGQL	117.000	enhance P2	SEQ ID NO:129
2	31	EPCGFEATYL	4.800		
3	66	EIEINGQLVF	3.000		
4	26	VVEYCEPCGF	3.000		
5	50	YPGIEIESRL	2.600		
HLA-	-B*3901				
1	94	RRASNGETL	15.000		
exam _l impro	ple of wed peptide	RHASNGETL	90.000	enhance P2	SEQ ID NO:130
2	34	GFEATYLEL	9.000		
3	38	TYLELASAV	4.000		
4	66	EIEINGQLV	3.000		
5	2	SGEPGQTSV	3.000		
HLA-	-B*3901 (10-	-mer peptides)			
1	33	CGFEATYLEL	12.000		
exam _l impro	ple of eved peptide	CHFEATYLEL	360.000	enhance P2	SEQ ID NO:131
2	64	AFEIEINGQL	9.000		
3	93	IRRASNGETL	4.500		
4	46	VKEQYPGIEI	3.000		·
5	16	EVEPGSGVRI	3.000		

	Examples of predicted human Class I MHC binding peptides - c ntinued Rank Start Subsequence Score								
Kank	Position	Subsequence	Score (estimated half time of dissociation)		SEQ ID NO.				
	D+2002								
nla	-B*3902								
1	70	NGQLVFSKL	2.400						
exam _i	ple of oved peptide	NKQLVFSKL	24.000	enhance P2	SEQ ID NO:132				
2	81	GGFPYEKDL	2.400						
3	94	RRASNGETL	2.000						
4	34	GFEATYLEL	2.000						
5	107	NSRPPCVIL	0.600						
HLA	-B*3902 (10-	mer peptides)							
1	69	INGQLVFSKL	2.400						
2	64	AFEIEINGQL	2.400						
3	50	YPGIEIESRL	2.400		,				
4	80	NGGFPYEKDL	2.400						
5	106	TNSRPPCVIL	2.000						
HLA	-B*4403								
1	67	IEINGQLVF	200.000						
	ple of oved peptide	IEINGQLVY	900.000	enhance P9	SEQ ID NO:133				
2	27	VEYCEPCGF	40.000						
3	21	SGVRIVVEY	36.000						
4	65	FEIEINGQL	20.000		•				
5	35	FEATYLELA	12.000						

	ples of predic Start Position	cted human Class Subsequence	Score	ng peptides – continued half time of dissociation)	SEQ ID NO.
HLA	-B*4403 (10-	mer peptides)			
1	30	CEPCGFEATY	120.000		
2	53	IEIESRLGGT	30.000		
	ple of oved peptide	IEIESRLGGY	900.000	enhance P10	SEQ ID NO:134
3 .	67	IEINGQLVFS	30.000		
4	65	FEIEINGQLV	20.000		
5	17	VEPGSGVRIV	18.000		
HLA	-B*5101				
1	18	EPGSGVRIV	484.000		
2	59	LGGTGAFEI	114.400		
	ple of oved peptide	LPGTGAFEI	572.000	enhance P2	SEQ ID NO:135
3	2	SGEPGQTSV	48.400		
4	81	GGFPYEKDL	44.000		
5	70	NGQLVFSKL	22.000		
HLA	B*5101 (10-	mer peptides)			
1	18	EPGSGVRIVV	440.000		
2	44	SAVKEQYPGI	220.000		
	ple of oved peptide	SPVKEQYPGI	440.000	enhance P2	SEQ ID NO:136
3	31	EPCGFEATYL	220.000		
4	81	GGFPYEKDLI	176.000		
5	50	YPGIEIESRL	157.300		

	ples of predic Start Position	cted human Class I Subsequence	Score	ng peptides – continued alf time of dissociation)	SEQ ID NO.
HLA	-B*5102				
1	18	EPGSGVRIV	242.000		
2	81	GGFPYEKDL	110.000		
	ple of oved peptide	GP FPYEKDI	2200.000	enhance P2, P9	SEQ ID NO:137
3	59	LGGTGAFEI	96.800		
4	70	NGQLVFSKL	48.400		
5	2	SGEPGQTSV	24.200		
HLA	-B*5102 (10-	-mer peptide)			
1	44	SAVKEQYPGI	726.000		
	ple of wed peptide	SPVKEQYPGI	1452.000	enhance P2	SEQ ID NO:138
2	50	YPGIEIESRL	400.000		
3	81	GGFPYEKDLI	400.000		
4	18	EPGSGVRIVV	220.000		
5	31	EPCGFEATYL	121.000		
HLA	-B*5103				
1	59	LGGTGAFEI	48.400		,
	ple of oved peptide	LAFTGAFEI	145.200	enhance P2	SEQ ID NO:139
2	2	SGEPGQTSV	44.000		
3	18	EPGSGVRIV	44.000		
4	70	NGQLVFSKL	7.260		
5	81	GGFPYEKDL	7.200		

Exam	nles of predic	cted human Class l	MHC hindir	ng peptides – continued	Appl. No. 09/824,/8/
	Start	Subsequence	Score		
	Position		(estimated h	alf time of dissociation)	SEQ ID NO
HLA	-B*5103 (10-	-mer peptide)			
1	44	SAVKEQYPGI	110.000		
2	81	GGFPYEKDLI	52.800		
3	18	EPGSGVRIVV	44.000		
	ple of wed peptide	EAGSGVRIVV	110.000	enhance P2	SEQ ID NO:140
4	60	GGTGAFEIEI	44.000		
5	33	CGFEATYLEL	7.920		
HLA-	-B*5201				
1	18	WPGSGVRIV	75.000		
2	67	LEINGQLVF	22.500		
	ple of wed peptide	L Q INGQLV I	450.000	enhance P2, P9	SEQ ID NO:141
3	59	LGGTGAFEI	11.250		
4	98	NGETLEKIT	11.000		
5	19	PGSGVRIVV	10.000		
HLA:	-B*5201 (10-	-mer peptides)			
1	18	EPGSGVRIVV	100.000		
2	17	VEPGSGVRIV	45.000		
	ple of oved peptide	V Q PGSGVRIV	450.000	enhance P2	SEQ ID NO:142
3	81	GGFPYEKDLI	33.000		
4	105	ITNSRPPCVI	15.000		
5	37	ATYLELASAV	12.000		

	ples of predic	cted human Class I Subsequence	MHC bindin	ng peptides – continued	Appi. No. 09/824,/8/
	Position			alf time of dissociation)	SEQ ID NO.
HLA:	-B*5801				
1	75	FSKLENGGF	40.000		
exam _l	ple of wed peptide	FSKLENGGW	80.000	enhance P9	SEQ ID NO:143
2	42	LASAVKEQY	4.500		
3	107	NSRPPCVIL	4.000		
4	61	GTGAFEIEI	3.000		
5	105	ITNSRPPCV	3.000		
HLA	-B*5801 (10-	-mer peptides)			
1	56	ESRLGGTGAF	12.000		
2	20	GSGVRIVVEY	10.800		
	ple of wed peptide	GSGVRIVVEW	144.000	enhance P10	SEQ ID NO:144
3	1	MSGEPGQTSV	4.000		
4	105	ITNSRPPCVI	3.000		٠
5	37	ATYLELASAV	3.000		
HLA	-Cw*0301				
1	65	FEIEINGQL	30.000		
2	81	GGFPYEKDL	18.000		
3	70	NGQLVFSKL	12.000		
4	57	SRLGGTGAF	10.000		
5	34	GFEATYLEL	10.000		

Examples of predicted human Class I MHC binding peptides - continued

Ranl	Start Position	Subsequence	Score (estimated)	half time of dissociation)	SEQ ID NO.
HLA	A-Cw*0301 (1	0-mer peptides)			
1	44	SAVKEQYPGI	50.000		
	nple of oved peptide	SAVKEQYPGL	100.000	enhance P10	SEQ ID NO:145
2	33	CGFEATYLEL	45.000		
3	69	INGQLVFSKL	12.000		
4	81	GGFPYEKDLI	3.750		
5	106	TNSRPPCVIL	3.000		
HL	A-Cw*0401				
1	34	GFEATYLEL	240.000		
2	38	TYLELASAV	30.000		
3	82	GFPYEKDLI	25.000		
4	18	EPGSGVRIV	20.000		
5	31	EPCGFEATY	12.000		
	nple of roved peptide	EFCGFEATL	200.000	enhance P2, P9	SEQ ID NO:146
HL	A- Cw*0401 (1	0-mer peptides)			
1	64	AFEIEINGQL	200.000		
2	74	VFSKLENGGF	100.000		
	mple of roved peptide	VFSKLENGGL	200.000	enhance P10	SEQ ID NO:147
3	50	YPGIEIESRL	80.000		
4	31	EPCGFEATYL	80.000		
5	18	EPGSGVRIVV	10.000		

Exan Rank	nples of pred Start Position	licted human Class Subsequence	I MHC binding peptides – continued Score (estimated half time of dissociation)	SEQ ID NO.
HLA	-Cw*0602			
1	85	YEKDLIEAI	6.600	
2	65	FEIEINGQL	6.600	
3	21	SGVRIVVEY	6.000	
4	31	EPCGFEATY	3.300	
5	61	GTGAGEIEI	3.000	
HLA.	-Cw*0702			
1	31	EPCGFEATY	24.000	
2	21	SGVRIVVEY	19.200	•
3	42	LASAVKEQY	8.800	
1	77	KLENGGFPY	4.000	
5	49	QYPGIEIES	2.880	
ILA-	Cw*0702 (1	10-mer peptides)		
	20	GSGVRIVVEY	38.400	
!	30	CEPCGFEATY	16.000	
	41	ELASAVKEQY	16.000	
	50	YPGIEIESRL	7.920	
	76	SKLENGGFPY	4.000	

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